

Edgar Filing: Ultragenyx Pharmaceutical Inc. - Form 8-K

Ultragenyx Pharmaceutical Inc.  
Form 8-K  
April 17, 2018  
UNITED STATES  
SECURITIES AND EXCHANGE COMMISSION  
WASHINGTON, DC 20549

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FORM 8-K

CURRENT REPORT

Pursuant to Section 13 or 15(d) of the  
Securities Exchange Act of 1934

Date of Report (Date of earliest event reported): April 17, 2018

ULTRAGENYX PHARMACEUTICAL INC.

(Exact name of registrant as specified in charter)

Delaware	001-36276	27-2546083
(State or other jurisdiction of incorporation)	(Commission (IRS Employer File Number)	Identification No.)

60 Leveroni Court, Novato, California 94949  
(Address of principal executive offices) (Zip Code)

Registrant's telephone number, including area code: (415) 483-8800

Not Applicable

(Former name or former address, if changed since last report)

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:

Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)

Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)

Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))

Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§ 230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§ 240.12b-2 of this chapter).

Emerging growth company

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If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act.

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Item 8.01. Other Events.

On April 17, 2018, Ultragenyx Pharmaceutical Inc. (the “Company”) issued a press release (the “Release”) announcing that the U.S. Food and Drug Administration has approved Crysvita® (burosumab-twza) for the treatment of X-linked hypophosphatemia (XLH) in adult and pediatric patients one year of age and older.

A copy of the Release is filed herewith as Exhibit 99.1.

Item 9.01 Financial Statements and Exhibits

(d) Exhibits

Exhibit No. Description

99.1 Press Release of Ultragenyx Pharmaceutical Inc., dated April 17, 2018

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SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

Date: April 17, 2018      Ultragenyx Pharmaceutical Inc.

By: /s/ Shalini Sharp  
Shalini Sharp  
Executive Vice President, Chief Financial Officer